

LPAAT- β Inhibitors and Uses Thereof

Cross-reference Information

5 This application is a continuation of U.S. Patent Application Serial No. 09/984,888 filed October 31, 2001 which claims priority to U.S. Provisional Application Serial No. 60/244,195, filed October 31, 2000, the disclosures of which are incorporated by reference herein in their entirety.

10 *Background of the Invention*

Field of the Invention

 The invention is in the field of organic and medicinal chemistry. In particular, the invention relates to triazines and the use thereof to inhibit
15 lysophosphatidic acid acyltransferase β (LPAAT- β) activity. The invention further relates to methods of treating cancer using said triazines. The invention also relates to methods for screening for LPAAT- β activity.

Related Art

20 LPAAT catalyzes the acylation of lysophosphatidic acid (LPA) to phosphatidic acid. LPA is the simplest glycerophospholipid, consisting of a glycerol molecule, a phosphate group, and a fatty acyl chain. LPAAT adds a second fatty acyl chain to LPA, producing phosphatidic acid (PA). PA is the precursor molecule for certain phosphoglycerides, such as phosphatidylinositol,
25 and diacylglycerols, which are necessary for the production of other phosphoglycerides, such as phosphatidylcholine, and for triacylglycerols, which are essential biological fuel molecules.

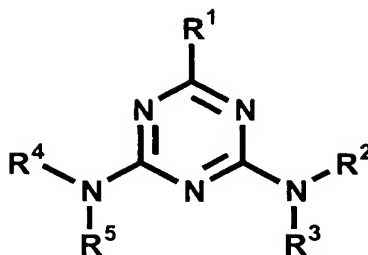
 In addition to being a crucial precursor molecule in biosynthetic reactions, LPA has recently been added to the list of intercellular lipid messenger
30 molecules. LPA interacts with G protein-coupled receptors, coupling to various independent effector pathways including inhibition of adenylate cyclase, stimulation of phospholipase C, activation of MAP kinases, and activation of the small GTP-binding proteins Ras and Rho. Moolenaar, *J. Biol. Chem* 28:1294 (1995). The physiological effects of LPA have not been fully characterized as

yet. However, one of the physiological effects that is known is that LPA promotes the growth and invasion of tumor cells. It has been shown that the addition of LPA to ovarian or breast cancer cell lines induces cell proliferation, increases intracellular calcium levels, and activates MAP kinase. Xu *et al.*,
5 *Biochem. J.* 309:933 (1995). In addition, LPA has been shown to induce MM1 tumor cells to invade cultured mesothelial cell monolayers. Imamura *et al.* *Biochem. Biophys. Res. Comm.* 193:497 (1993).

Like LPA, PA is also a messenger molecule. PA is a key messenger in a common signaling pathway activated by proinflammatory mediators such as
10 interleukin-1 β , tumor necrosis factor α , platelet activating factor, and lipid A. Bursten *et al.*, *Am. J. Physiol.* 262:C328 (1992); Bursten *et al.*, *J. Biol. Chem.* 255:20732 (1991); Kester *J. Cell Physiol.* 156:317 (1993). PA has been implicated in mitogenesis of several cell lines [English, *Cell Signal* 8: 341 (1996)]. PA level has been found to be increased in either ras or fps transformed
15 cell lines compared to the parental Rat2 fibroblast cell line [Martin *et al.*, *Oncogene* 14: 1571 (1997)]. Activation of Raf-1, an essential component of the MAPK signaling cascade, by extracellular signals is initiated by association with intracellular membranes. Recruitment of Raf-1 to membranes has been reported to be mediated by direct association with phosphatidic acid [Rizzo *et al.*, *J Biol*
20 *Chem* 275:23911-8 (2000)]. Thus, LPAAT, as an enzyme that regulate PA content in cells, may play a role in cancer, and may also mediate inflammatory responses to various proinflammatory agents.

Summary of the Invention

25 The preferred embodiments of the present invention relate to a compound of the Formula:



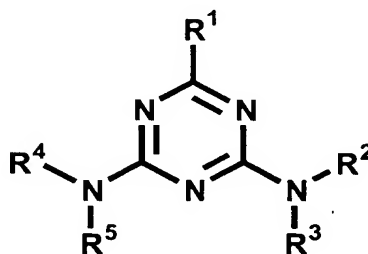
wherein,

R¹ is halo, hydroxy, alkylmercapto, mercapto, alkoxy, aryloxy or
5 substituted amino;

R², R³, R⁴ and R⁵, each of which may be same or different, are hydrogen,
alkyl, substituted alkyl, alkenyl, alkynyl, aryl or substituted aryl; or

R² and R³ or R⁴ and R⁵, together with the nitrogen to which they are
attached, form a piperidine, piperazine, or a morpholine ring; or
10 pharmaceutically acceptable salts thereof.

The preferred embodiments of the present invention further relate to a
method for inhibiting LPAAT-β (lysophosphatidic acid acyltransferase β)
comprising contacting LPAAT-β with an effective amount of a compound of the
Formula:



15

wherein,

R¹ is halo, hydroxy, alkylmercapto, mercapto, alkoxy, aryloxy or
substituted amino;

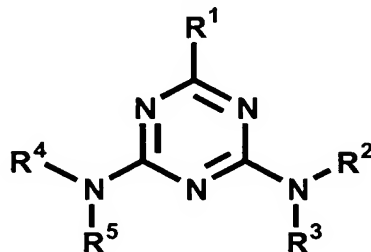
R², R³, R⁴ and R⁵, each of which may be same or different, are hydrogen,
20 alkyl, substituted alkyl, alkenyl, alkynyl, aryl or substituted aryl; or

R² and R³ or R⁴ and R⁵, together with the nitrogen to which they are
attached, form a piperidine, piperazine, or a morpholine ring; or

pharmaceutically acceptable salts thereof;

thereby inhibiting LPAAT-β.

The preferred embodiments of the present invention further relate to a method of inhibiting cell proliferation comprising contacting a cell with an effective amount of a compound of the Formula:



5 wherein,

R¹ is halo, hydroxy, alkylmercapto, mercapto, alkoxy, arylox or substituted amino;

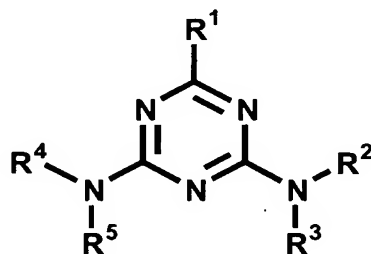
R², R³, R⁴ and R⁵, each of which may be same or different, are hydrogen, alkyl, substituted alkyl, alkenyl, alkynyl, aryl or substituted aryl; or

10 R² and R³ or R⁴ and R⁵, together with the nitrogen to which they are attached, form a piperidine, piperazine, or a morpholine ring; or

pharmaceutically acceptable salts thereof;

thereby inhibiting the proliferation of the cell.

The preferred embodiments of the present invention further relate to a
15 method for treating cancer, comprising administering to an animal in need thereof, an effective amount of a compound of the Formula:



wherein,

20 R¹ is halo, hydroxy, alkylmercapto, mercapto, alkoxy, aryloxy or substituted amino;

R², R³, R⁴ and R⁵, each of which may be same or different, are hydrogen, alkyl, substituted alkyl, alkenyl, alkynyl, aryl or substituted aryl; or

R² and R³ or R⁴ and R⁵, together with the nitrogen to which they are attached, form a piperidine, piperazine, or a morpholine ring; or

25

pharmaceutically acceptable salts thereof;
wherein the cancer is treated.

The preferred embodiments of the present invention further relate to a method for screening a patient for LPAAT- β activity, said method comprising
5 detecting the presence or absence of an increased amount of LPAAT- β RNA, DNA or protein relative to a predetermined control, whereby the presence of said increased amount is indicative of cancer susceptibility in said patient.

The preferred embodiments of the present invention further relate to a method of inhibiting cell proliferation comprising the inhibition of LPAAT- β .

10 The preferred embodiments of the present invention further relate to a vaccine preparation capable of inducing an anti-tumor immune response comprising a pharmaceutically acceptable carrier and an anti-tumor immune response-inducing effective amount of LPAAT- β protein.

The preferred embodiments of the present invention further relate to a
15 method for screening a patient for LPAAT- β activity, said method comprising detecting the presence or absence of an increased amount of a phospholipid of defined acyl-chain composition relative to a predetermined control, whereby the presence of said increased amount is indicative of cancer susceptibility in said patient.

20

Detailed Description of the Preferred Embodiments

I. LPAAT- α and LPAAT- β : An Overview.

Northern blot analysis shows that LPAAT- α is expressed in all human
25 tissues tested with the highest expression level found in skeletal muscle (West *et al. DNA Cell Biol.* 16:691 (1997)). The uniformity of LPAAT- α expression has also been found in additional tissues such as prostate, testis, ovary, small intestine, and colon (Stamps *et al., Biochem. J.* 326: 455 (1997)) as well as in mouse tissues (Kume *et al., Biochem. Biophys. Res. Commun.* 237: 663 (1997)).
30 A 2 kb and a 1.3 kb forms, possibly due to alternative utilization of polyadenylation signals at the 3'-UTR, have been found in murine LPAAT- α mRNA (Kume *et al., Biochem. Biophys. Res. Commun.* 237: 663 (1997)), whereas

only one major human LPAAT- α mRNA of 2 kb in size has been detected by Northern analysis. West *et al.*, *DNA Cell Biol.* 16: 691 (1997); Stamps *et al.*, *Biochem. J.* 326: 455 (1997).

In contrast, LPAAT- β demonstrates a distinct tissue distribution of mRNA expression. West *et al.*, *DNA Cell Biol.* 16: 691 (1997). LPAAT- β is most highly expressed in liver and heart tissues. LPAAT- β is also expressed at moderate levels in pancreas, lung, skeletal muscle, kidney, spleen, and bone marrow; and at low levels in thymus, brain and placenta. This differential pattern of LPAAT- β expression has been confirmed independently (Eberhardt *et al.*, *J. Biol. Chem.* 272: 20299 (1997)) with the only discrepancy being that high level, instead of moderate level, of LPAAT- β has been detected in pancreas, possibly due to slight lot variations in commercial RNA blots (Clontech, Palo Alto, CA). In addition, moderate LPAAT- β expression has been found in prostate, testis, ovary, small intestine, and colon with the small intestine containing relatively higher amounts. Eberhardt *et al.*, *J Biol Chem* 272: 20299 (1997). Within various brain sections, high expression has been found in the subthalamic nucleus and spinal cord; and least in the cerebellum, caudate nucleus, corpus callosum, and hippocampus. LPAAT- β can also be detected in myeloid cell lines THP-1, HL-60, and U937 with the mRNA levels remaining the same with or without phorbol-ester treatment. The size difference between human LPAAT- α and LPAAT- β mRNA is consistent with the sequence data, in which LPAAT- α has a longer 3'-UTR. The differential tissue expression pattern LPAAT- α and LPAAT- β mRNA would suggest these two genes are regulated differently and are likely to have independent functions. Therefore, a desirable feature in compounds that inhibit LPAAT activity is that they are specific in inhibiting one isoform of the enzyme over the other (i.e., LPAAT- β over LPAAT- α).

II. LPAAT- β and Cancer.

PA has been implicated in mitogenesis of several cell lines. English, *Cell Signal* 8: 341 (1996). PA level has been found to be increased in either ras or fps transformed cell lines compared to the parental Rat2 fibroblast cell line (Martin *et al.*, *Oncogene* 14: 1571 (1997). To test whether LPAAT expression may be

enhanced in certain tumor cells, the expression of LPAAT- α and LPAAT- β mRNA in human tumor panel blots (Invitrogen, Carlsbad, CA) that contained tumor RNAs, isolated from various malignant tissues and RNAs from the normal tissues in the surgical margins, were examined. Leung *et al.*, *DNA Cell Biol.* 17: 377 (1998). The same blots were also reprobbed using cDNAs encoding phosphatidic acid phosphatase isoform PAP2- α ; an enzyme that degrades, rather than generates, PA. Of a total of eight different tissues examined, LPAAT- β mRNA was found to be elevated in three tumor tissues (uterus, fallopian tube, and ovary), as compared to its expression in the corresponding normal tissues. However, no significant difference was found in LPAAT- α mRNA level between the various tumor tissues and the normal adjacent tissues. In two of the tumor tissues (fallopian tube and ovary) where LPAAT- α mRNA was elevated, PAP2- α mRNA expression was found to be suppressed, as it was also in tumors of the colon, rectum, and breast.

Since the finding of differential expression of LPAAT- β mRNA in certain tumor versus normal tissues is based on Northern analysis of a single specimen from a given tissue, more studies will be needed to determine whether the relative elevation of LPAAT- β expression in selected tumor tissues can be applied and extended to similar tissues derived from a larger number of donors. Leung *et al.*, *DNA Cell Biol.* 17: 377 (1998). Accordingly, *in situ* hybridization was used to compare LPAAT- β mRNA levels in breast, ovary, and prostate tumor samples obtained from multiple independent donors (LifeSpan Biosciences, Seattle, WA). Specifically, the coding region of human LPAAT- β was amplified by PCR from the plasmid pCE9. LPAAT- β with primers 5'-GCATGAATTC AAAGGCCTAC GTCGACATGG AGCTGTGGCC GTG-3' and 5'-GTCGACTCTA GACTACTGGG CCGGCTGCAC-3'. The resultant 870 bp PCR product was then cut with EcoR I and XbaI for insertion in between the EcoR I and XbaI sites of the *in vitro* transcription vector pDP18-T7/T3 (Ambion, Austin, TX) to generate the plasmid pDP_lptB. Serial tissue sections from paraffin archival samples were hybridized with digoxigenin labeled riboprobes transcribed from either a T3 (sense) or T7 (antisense) transcription initiation site present in the plasmid pDP_lptB linearized with either EcoR I (antisense) or Xba I (sense). The

tissue sections from paraffin blocks were digested with proteinase K (20 µg/ml) for 4 minutes, then hybridized with the antisense probe (1 µg/ml) at 60° C for 22 hours and subsequently washed with 2xSSC and 0.1xSSC at 50° C. The hybridization signals were detected with NBT/BCIP substrates using three cycles
5 of an alkaline phosphatase TSA amplification system (NEN Life Sciences, Boston, MA). The specimens were then counterstained with methyl green. The signal was developed within 30 minutes at room temperature. The slides were then imaged using a digital camera mounted onto a microscope.

Breast and ovary tissues were chosen for further *in situ* hybridization
10 study, as initial Northern analysis showed elevation of LPAAT-β mRNA levels in tumors derived from the female reproductive tract. Prostate tissue was chosen, as it responds to steroid hormones and contains ductal structures in a manner similar to breast and ovary tissues. Using an anti-sense cDNA probe, it was demonstrated that expression of the β isoform of this enzyme (LPAAT-β) was
15 augmented in human tumor tissue in 10/11 ovarian, 14/20 breast, and 7/16 prostate biopsies as compared to normal adjacent tissues. Figure 1 shows an example of the results on a breast intraductal adenocarcinoma sample where there is moderate increase in LPAAT-β mRNA level in the tumor samples (top 2 panels) as evidenced by more dark-purple to brown spots compared to adjacent
20 hyperplasia (bottom-left panel) and normal tissue (bottom-right panel). The slight increase in LPAAT-β mRNA staining in the hyperplasia sample (bottom-left panel) versus the normal sample (bottom-right panel) suggests that elevation occurs at an early stage of oncogenesis. Figure 2 shows an example of the results on another breast intraductal adenocarcinoma sample where there is large increase
25 in LPAAT-β mRNA level in the tumor sample (left panel) as evidenced by more dark-purple spots versus the adjacent normal tissue (right panel). Figure 3 shows three examples of ovarian cancer samples where the LPAAT-β mRNA levels are elevated and one example with undetectable level of LPAAT-β mRNA (lower right panel). Figure 4A shows an example of the results on a prostate
30 adenocarcinoma sample where there is moderate increase in LPAAT-β mRNA level in the tumor samples (left panel) as evidenced by more dark-purple spots versus the adjacent normal tissue (right panel). In no cases have elevated levels

of LPAAT- β mRNA expression been found in the adjacent normal region from the same donor even in those cases of breast, ovarian, or prostate tumor where LPAAT- β mRNA levels happen to be low or undetectable. The augmented expression of LPAAT- β in a high percentage of tumor samples from breast
5 (70%), ovary (91%), and prostate tissues (44%) would suggest that LPAAT- β overexpression may be a contributing factor for the development of these tumors.

To determine if increased transcription of LPAAT- β mRNA in selected tumor tissues can be extended to increased LPAAT- β protein expression in a wider range of tissues, a monoclonal antibody specific for human LPAAT- β
10 protein (MoAb 4B12) was generated based on the peptide sequence, DLGERMVRENKLVW, derived from amino acids 155-168 of LPAAT- β protein (BAbCO, Berkeley, CA). Figure 4B shows an example of the results on immunohistochemical staining (PhenoPath, Seattle, WA) with MoAb 4B12 at 1:4000 dilution of ovarian tissue where there is substantial increase in LPAAT- β
15 protein expression in the tumor samples (right panels) as evidenced by more intense brown stainings versus the normal tissue (left panel). Figure 4C shows an example of the results on immunohistochemical staining (PhenoPath, Seattle, WA) with MoAb 4B12 at 1:4000 dilution of cervical tissue where there is substantial increase in LPAAT- β protein expression in the tumor samples (right
20 panels) as evidenced by more intense brown stainings versus the normal tissue (left panel). There is also more staining in the surrounding stromal cells (indicated by arrows) in the tumor tissue vs the normal tissue, suggesting that the tumor may also induce LPAAT- β protein expression in the surrounding cells. Figure 4D shows another example of the results on immunohistochemical
25 staining (PhenoPath, Seattle, WA) with MoAb 4B12 at 1:4000 dilution of lung tissue where there is extensive increase in LPAAT- β protein expression in the tumor samples (right panels) as evidenced by more intense brown stainings versus the normal tissue (left panel). Figure 4E shows the summary of immunohistochemistry (IHC) results of the various tissue samples stained by
30 MoAb 4B12. The augmented expression of LPAAT- β in a high percentage of tumor samples again suggest that LPAAT- β overexpression may be a contributing factor for the development of these tumors and that LPAAT- β may be a useful

target for the development of anti-cancer compounds.

The aforementioned antibody may also be used for diagnostic and prognostic purposes when a tumor is present both on biopsies and in serum or plasma. For example, ELISA may be performed on serum to detect lung or ovarian cancer. It should be mentioned that currently there are no useful early
5 diagnostics for these types of cancers.

The overexpression of LPAAT- β in selected tumor tissues would also suggest the LPAAT- β protein may constitute a useful antigen for the development of tumor vaccines against those tumors where LPAAT- β is overexpressed. Fong *et al.*, *Annu. Rev. Immunol.* 18: 245 (2000); Schreurs, *et al.*,
10 *Crit. Rev. Oncol.* 11: 1 (2000). One such approach may use autologous dendritic cells, a type of potent antigen-presenting cells, to present LPAAT- β as a tumor-associated antigens for the generation of tumor-specific immunity through the MHC class I and II processing pathways. Administration of dendritic cells
15 loaded *ex vivo* with LPAAT- β as a therapeutic vaccine to patients with tumors with augmented LPAAT- β expression may induce T cell-mediated tumor destruction.

To assess whether LPAAT- β overexpression in cells would lead to certain phenotypic changes that are commonly observed in transformed cells, the growth
20 and adherence characteristics of ECV304 cells (American Type Culture Collection, Richmond, VA) expressing LPAAT- β (LPTb), expressing a catalytically inactive form of LPAAT- β (b-M8) whereby the arginine at position 175 was changed to alanine using the GeneEditor™ *in vitro* site-directed mutagenesis system (Promega, Madison, WI), or expressing green fluorescent
25 protein (GFP) as a control were compared. The aforementioned cells that express GFP may be considered to be a non-limiting example of a “predetermined control,” according to the preferred embodiments of the present invention. That is, such cells may be used to gauge whether a cell is over- or under-expressing LPAAT- β DNA, RNA or protein. Figure 5A shows the growth curve of these
30 three cell lines. Each cell line was seeded at 200,000 cells per 60 mm plate. The cell numbers at various times after seeding were determined by counting with a hemacytometer. The growth rate of the three cell lines were similar until they reached confluence at 100 hours after plating. After confluence, the LPTb cells

were able to continue to proliferate, while the b-M8 and GFP cells' growth started to level off. This demonstrated that ECV304 cells overexpressing LPAAT- β could continue to grow and could form a plurality of layers after they had formed a confluent monolayer of cells. The proliferation of the cells with the inactive mutant or the control cells slowed down after confluence. The loss of contact inhibition and the propensity for growth to an unusually high cell density are changes commonly observed in tumorigenesis. The fact that the inactive LPAAT- β mutant (b-M8) expressing cells, like the vector control cells, are constrained by density-dependent inhibition of cell division strongly suggests that the capacity to overcome contact inhibition may be due to increases in LPAAT- β enzymatic activity. The development of compounds that inhibit LPAAT- β enzymatic activity may reverse the growth pattern and hence tumorigenesis in cells with abnormally high level of LPAAT- β expression.

To determine if the observation from LPAAT- β expressing ECV304 cells can be extended to other cell types and to animal models of tumorigenesis, LPAAT- β cDNA was inserted into a retroviral expression vector, pLOXSN, for the generation of recombinant viral stocks in a packaging cell line, PT67 (Clontech, Palo Alto, CA), for transduction into various cell lines. The vector pLOXSN was derived from pLXSN with insertion of a 19bp oligonucleotide coding for the locus of recombination (lox) signal sequence as well as a ClaI recognition site into the NheI site within the 3'-LTR region of pLXSN. Miller and Rosman *BioTechniques* 7: 980 (1989); Hoess. and Abremski, *Nucleic Acid and Mol. Biol.* 4: 99 (1990). This lox sequence will be duplicated within the 5'-LTR region during viral replication. Hence the sequence in between the two lox sites located within the 5'- and the 3'-LTR can be excised if required in the presence of the enzyme cre recombinase supplied *in trans* from a separate retroviral vector with a different selectable marker.

Over-expression of the normal cellular LPAAT- β cDNA in NIH/3T3 cells was associated with transformation in 3 out of 9 transduced populations. As is the case with normal cellular proto-oncogenes, over-expression of LPAAT- β is not sufficient, but may contribute to transformation along with other, spontaneous events. Figure 5B shows examples of cell morphology of NIH/3T3 cells: a bulk

population transfected with a plasmid overexpressing the Ki-ras oncogene (top left panel), a selected clone transduced with a retroviral vector overexpressing LPAAT- β (Hc2; lower left panel) and cells with the LPAAT- β cDNA excised using the lox-cre recombination in the lower left and normal, untransduced cells (top right panel). Sauer, *Methods* 14: 381 (1998). The control untransduced cells exhibited normal fibroblast morphology and grew as a contact-inhibited, adherent monolayer (top right panel). In contrast, both the Ki-ras and LPAAT- β overexpressing cells were more elongated and spiked, were not contact-inhibited and formed foci typical of transformation of these immortalized fibroblasts. After removal LPAAT- β transgene by lox-cre recombination from the Hc2 clone (bottom right panel), this transformed morphology was lost, suggesting that LPAAT- β overexpression is a contributing factor to this transformation phenotype rather than being the result entirely of spontaneous events during in vitro passage.

Another common parameter of cancer cells is a reduced requirement for elements present in serum. Figure 5C compares the growth profiles of transduced populations of NIH/3T3 cells in low (2%) serum. Two independent populations (LPT Hc2, LPT L bulk) overexpressing LPAAT- β have an increased ability to proliferate compared to a control vector clone expressing alkaline phosphatase (APc1) and those corresponding populations with deletion of the LPAAT- β transgene by lox-cre recombination (LPT Hc2cre, LPT L bulkcre), suggesting that LPAAT- β overexpression is a contributing factor to this transformed phenotype of proliferation with a reduced requirement for growth factors.

Similarly, out of a total of 12 populations of human prostate LNCaP cells (American Type Culture Collection, Manassas, VA) transduced with LPAAT- β expressing vector, most of them show augmented proliferation in low serum when compared to control cells (Figure 5D).

To determine whether administration of LPAAT- β inhibitor would have any effect on cell proliferation in tissue culture, proliferation of human breast tumor MCF-7 cells in microplates were measured by CyQUANT analysis using a green-fluorescent nucleic acid stain optimized to produce a linear detection range from 50 to 50,000 cells in 200 μ l volume (Molecular Probes, Eugene, OR) in the

presence of various concentrations of a LPAAT- β inhibitor. Figure 5E shows the triazine compound shows 6-chloro-N,N'-diphenyl-[1,3,5]triazine-2,4-diamine at $\geq 20 \mu\text{M}$ is effective in blocking the proliferation of MCF-7 cells.

To determine if LPAAT- β overexpression would contribute to tumorigenesis in mice, 2×10^6 NIH3T3 cells overexpressing LPAAT- β (LPAAT vector) and control cells were injected subcutaneously into nude mice. Figure 6A shows tumor could be detected after 14 days from the LPAAT- β overexpressing cells, while no tumor formation was detected in vector control cells after 28 days. The cells with the transgene removed by lox-cre recombination showed delay of tumor formation compared to LPAAT- β overexpressing cells by ~ 7 days. Recovery and analysis of the lox-cre cells from mice showed that there had been *in vivo* selection of a small sub-population that had not been recombined to remove the LPAAT- β transgene. This analysis demonstrated that the only cells to form tumors retained the original LPAAT vector and indeed had a high level of LPAAT activity as well as G418 resistance (the neo gene is also removed along with LPAAT- β during the cre-lox procedure). These data show LPAAT- β overexpression is a contributing factor for tumorigenesis *in vivo*.

To determine whether administration of LPAAT- β inhibitor would have any effect on tumor growth in mice, 5×10^5 NIH/3T3 cells overexpressing the oncogene Ki-ras were injected subcutaneously into nude mice. An LPAAT- β inhibitor 6-chloro-N,N'-diphenyl-[1,3,5]triazine-2,4-diamine, at concentrations that range from 10 mg/Kg to 100 mg/Kg of mouse body weight was injected intra-peritoneally on day 1, 2, 3 and 4 after injection of tumor cells. The size of tumors was then measured on day 8. Figure 6B shows the volume of the tumors in mice is decreased as the concentration of the LPAAT- β inhibitor increases, suggesting that administration of this LPAAT- β inhibitor is efficacious in slowing down tumor growth *in vivo*.

In addition to slowing down tumor growth of NIH3T3/Ki-ras cells in nude mice, 6-chloro-N,N'-diphenyl-[1,3,5]triazine-2,4-diamine has also been found to decrease the growth of B16 melanoma cells (Figure 6C) and Lewis Lung tumor cells (Figure 6D) in syngeneic mice as well as the growth of human DU145 prostate tumor cells in nude mice in a xenograft study (Figure 6E).

Analysis and characterization of phospholipids and other complex lipids represent another strategy to measure effects of small molecule inhibitors on phospholipid metabolizing enzymes involved in tumor progression, including but not limited to, LPAAT- β . Measurements of phospholipids and other complex lipids may be derived from cell lines cultured *in vitro*, from tissue or plasma in vivo (e.g., murine or other animal studies), or from human subjects (e.g., phlebotomy or biopsy). Phospholipids, which are the primary constituents of a cellular bilayer, contain a universal phosphoric acid residue connected to a glycerol backbone. Phospholipid classes are defined by the chemical identity of the “head group” on the phosphoric acid moiety. However, each phospholipid class is often a complex mixture of discrete molecular species due to the fact that the glycerol backbone has two substituents residing at the Sn1 and Sn2 position of attachment. The substituents are acyl chains and typically consist of long chain fatty acids but may also include a long chain ether, acetyl, or hydroxyl group. Chemical measurements of phospholipids can involve a variety of analytical methods including, but not limited to, HPLC-MS (High Performance Liquid Chromatography-Mass Spectrometry), HPLC-MS/MS (High Performance Liquid Chromatography-Tandem Mass Spectrometry), one or two dimensional TLC (Thin Layer Chromatography), and radiometry. While all the stated methods can be used to quantitate bulk mass changes in a particular phospholipid class of interest, mass spectrometry offers the unique ability to measure all molecular species within a phospholipid class in a single measurement with a high degree of precision.

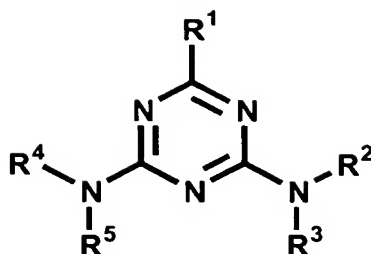
The above approach is demonstrated by performing HPLC-MS analyses of phospholipid extracts from murine NIH/3T3 immortalized fibroblasts, both normal wild type, β Hc2 cells (i.e., overexpressing LPAAT- β , and Hc2cre cells (i.e., LPAAT- β gene removed by site-specific recombination). Analysis of phosphatidylinositol in these cell populations clearly indicate a combined effect of LPAAT- β overexpression and cellular transformation for the Hc2 population over that of the normal wild type. This effect is characterized by an increase in unsaturated (i.e., palmitate and stearate) and monounsaturated (i.e., oleate) fatty acyl chains indicated by an increased molecular abundance of ions at m/z 807,

833, 835, 861, and 863 which correspond most likely to phosphatidylinositol species with acyl chains designated as 16:0-16:1, 16:1-18:1 (and/or 16:0-18:2), 16:0-18:1, 18:1-18:1 (and/or 18:0-18:2), and 18:0-18:1, respectively. While multiple molecular species may reside at the same nominal mass, these species
5 can be differentiated by tandem (MS/MS) mass spectrometry methods. Additionally, note that actual determination of positional location (i.e., Sn1 versus Sn2) requires other analytical methods and only the most prevalent configuration is listed here. In addition to the increase in unsaturated and monounsaturated acyl chains in the LPAAT- β overexpressing population (β Hc2),
10 there is also a corresponding decrease in polyunsaturated (i.e., arachidonate) fatty acyl chains at m/z 857 (16:0-20:4) and m/z 885 (18:0-20:4). Removal of the LPAAT- β transgene results in phosphatidylinositol distributions similar to that of the normal wild type 3T3 cells.

In summary, endogenous LPAAT- β expression is detected at high levels
15 by both *in situ* hybridization and immunohistochemistry in particular tumor tissues and often in surrounding stroma and is associated with tumor progression. LPAAT- β overexpression appears to contribute reversibly to transformation and tumorigenesis of immortalized rodent cells and may also contribute to increased transformation of weakly tumorigenic human cell lines. Compounds selected
20 from screening of LPAAT- β inhibitors from different structural families can inhibit proliferation of numerous tumor cell lines *in vitro*. Both nude and immunocompetent mice can tolerate at least 100 mg/kg/day for 4-5 days of the 6-chloro-N,N'-diphenyl-[1,3,5]triazine-2,4-diamine maintaining body weight and overall health with no discernable gross pathology. This compound can inhibit
25 the growth of numerous tumor models in mice and may be a tumor-static compound.

III. LPAAT- β Inhibitors.

In one aspect, the compounds of the present invention relate to triazines of
30 the Formula:



wherein,

R¹ is halo, hydroxy, alkylmercapto, mercapto, alkoxy, aryloxy or substituted amino;

5 R², R³, R⁴ and R⁵, each of which may be same or different, are hydrogen, alkyl, substituted alkyl, alkenyl, alkynyl, aryl or substituted aryl; or

R² and R³ or R⁴ and R⁵, together with the nitrogen to which they are attached, form a piperidine, piperazine, or a morpholine ring; or pharmaceutically acceptable salts thereof.

10 As used herein, "alkyl" refers to straight- or branched-chain hydrocarbons having from 1 to 10 carbon atoms and more preferably 1 to 8 carbon atoms which includes, by way of example, methyl, ethyl, *n*-propyl, *i*-propyl, *n*-butyl, *t*-butyl and the like.

The term "alkyl" also refers to an "unsaturated alkyl" moiety, which
15 means that it contains at least one alkene or alkyne moiety. "Alkene" or "alkenyl" refers to a group consisting of at least two carbon atoms and at least one carbon-carbon double bond. "Alkyne" or "alkynyl" refers to a group consisting of at least two carbon atoms and at least one carbon-carbon triple bond. The alkyl moiety, whether saturated or unsaturated, may be branched,
20 non-branched, or cyclic.

"Substituted alkyl" refers to an alkyl group, preferably containing from 1 to 10 carbon atoms, having from 1 to 5 substituents including halogen, hydroxyl, alkyl, aryl or substituted amino. A preferred substituted alkyl group is trifluoromethyl.

25 "Alkoxy" refers to the group "alkylBOB" which includes, by way of example, methoxy, ethoxy, *n*-propoxy, *i*-propoxy, *n*-butoxy, *t*-butoxy and the like.

"Substituted amino" refers to the group BNRR, wherein each R group is independently selected from the group consisting of hydrogen, alkyl, substituted

alkyl, cycloalkyl, aryl, substituted aryl, or the R groups can be joined together with the nitrogen to form a heterocyclic ring (e.g., piperidine, piperazine, or a morpholine ring).

"Aryl" refers to an unsaturated aromatic carbocyclic group of 6 to 14 carbon atoms having a single ring (e.g., phenyl) or multiple condensed rings (e.g., naphthyl or anthryl).

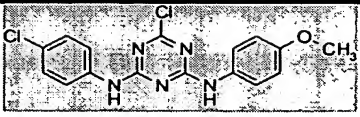
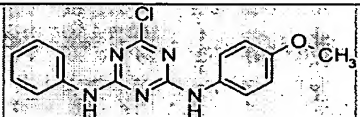
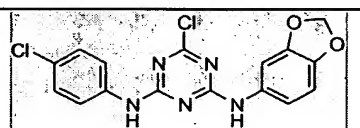
"Substituted aryl" refers to aryl group which are substituted with 1 to 3 substituents selected from hydroxy, alkyl, substituted alkyl, alkoxy, amino, aryl, $\text{O}-(\text{CH}_2)_n-\text{O}-$ (wherein n is an integer from 1 to 3), $-(\text{CH}_2)_m-$ (wherein m is an integer from 3 to 5) or halogen.

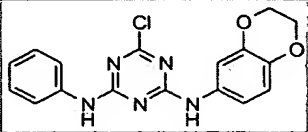
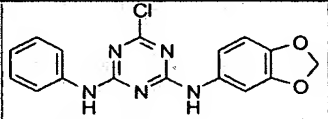
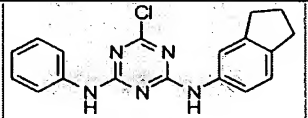
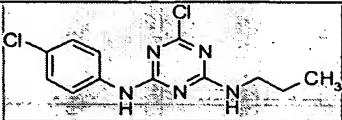
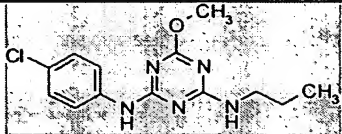
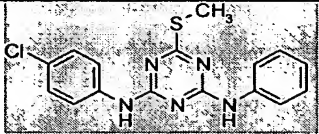
"Cycloalkyl" refers to cyclic alkyl groups containing between 3 and 8 carbon atoms having a single cyclic ring including, by way of example, cyclopropyl, cyclobutyl, cyclopentyl, cyclooctyl and the like.

"Halogen" or "halo" refers to fluoro, chloro, bromo, iodo. Most preferred halogens are chloro and fluoro.

"Mercapto" refers to the group $-\text{SR}$ wherein the R group is independently selected from the group consisting of hydrogen, alkyl, substituted alkyl, cycloalkyl, aryl or substituted aryl. The term "alkylmercaptop" refers to the group $-\text{SR}$ when R is alkyl, substituted alkyl or cycloalkyl.

Compounds of the preferred embodiments of the present invention include those compounds in Table 1.

CT Number	Structure	LPAAT β Colorimetric Assay IC_{50} (nM)
31867		750
31942		400
31978		1,000

32028		200
32042		200
32099		650
116988		160
117147		3,800
31888		3,100

Preferred compounds include, but are not limited to, 6-chloro-N-(4-methoxy-phenyl)-N'-p-tolyl-[1,3,5]triazine-2,4-diamine, N-butyl-6-chloro-N'-(4-chlorophenyl)-[1,3,5]triazine-2,4-diamine, 6-chloro-N-isopropyl-N'-p-tolyl-[1,3,5]triazine-2,4-diamine, N-tert-butyl-6-chloro-N'-phenyl-[1,3,5]triazine-2,4-diamine, (4-chloro-6-morpholin-4-yl-[1,3,5]triazin-2-yl)-naphthalen-1-yl-amine, N-tert-butyl-6-chloro-N'-p-tolyl-[1,3,5]triazine-2,4-diamine, 6-chloro-N-cyclohexyl-N'-isopropyl-[1,3,5]triazine-2,4-diamine, 2-(4-chloro-6-phenylamino-[1,3,5]triazin-2-ylamino)-2-methyl-propan-1-ol, 6-chloro-N-isopropyl-N'-phenyl-[1,3,5]triazine-2,4-diamine, 6-chloro-N-(4-chloro-phenyl)-N'-cyclohexyl-[1,3,5]triazine-2,4-diamine, N-allyl-6-chloro-N'-cyclohexyl-[1,3,5]triazine-2,4-diamine, 2-(4-chloro-6-phenylamino-[1,3,5]triazin-2-ylamino)-ethanol, N-tert-butyl-6-chloro-N'-cyclopentyl-[1,3,5]triazine-2,4-diamine, 6-chloro-N-(4-methoxyphenyl)-N'-phenyl-[1,3,5]triazine-2,4-diamine, N-benzo[1,3]dioxol-5-yl-6-chloro-N'-(4-chlorophenyl)-[1,3,5]triazine-2,4-diamine, 6-chloro-N-(2,3-dihydrobenzo[1,4]dioxin-6-yl)-N'-phenyl-[1,3,5]triazine-2,4-diamine, N-

benzo[1,3]dioxol-5-yl-6-chloro-N'-phenyl-[1,3,5]triazine-2,4-diamine, 6-chloro-N-indan-5-yl-N'-phenyl-[1,3,5]triazine-2,4-diamine, 6-chloro-N-(4-chlorophenyl)-N'-propyl-[1,3,5]triazine-2,4-diamine, N-(4-chlorophenyl)-6-methoxy-N'-propyl-[1,3,5]triazine-2,4-diamine and N-(4-chlorophenyl)-6-methylsulfanyln'
5 N'-phenyl-[1,3,5]triazine-2,4-diamine.

Most preferred compounds include, but are not limited to, 6-chloro-N,N'-diphenyl-[1,3,5]triazine-2,4-diamine, N-tert-butyl-6-chloro-N'-(4-chlorophenyl)-[1,3,5]triazine-2,4-diamine, 6-chloro-N-(4-chlorophenyl)-N'-(4-methoxyphenyl)-[1,3,5]triazine-2,4-diamine, 6-chloro-N-(4-chlorophenyl)-N'-phenyl-[1,3,5]-
10 triazine-2,4-diamine.

The compounds of the preferred embodiments of the present invention inhibit LPAAT- β and thereby inhibit cell proliferation. Therefore, the compounds of the preferred embodiments of the present invention may be useful in the treatment of cancer. The types of cancer that may be treated with the
15 compounds of the preferred embodiments of the present invention include, but are not limited to, prostate, breast, lung, ovarian, brain, cervical, colon or bladder cancer, and not limited to tumor cells expressing high levels of LPAAT- β as evidenced by the decrease in NIH/3T3 Ki-ras tumor cell growth *in vitro* and *in vivo* when treated with , 6-chloro-N,N'-diphenyl-[1,3,5]triazine-2,4-diamine.

20 **IV. Pharmacological Compositions, Therapeutic and Other Applications.**

The compound of the present invention, or its pharmaceutically acceptable salt, can be administered to a human patient *per se*, or in pharmacological compositions where it is mixed with pharmaceutically
25 acceptable carriers or excipient(s). Techniques for formulation and administration of drugs may be found in "Remington's Pharmaceutical Sciences," Mack Publishing Co., Easton, Pa., latest edition.

A. Routes Of Administration.

30 Suitable routes of administration may include, without limitation, oral, rectal, transmucosal or intestinal administration or intramuscular, subcutaneous,

intramedullary, intrathecal, direct intraventricular, intravenous, intraperitoneal or intranasal injections.

Alternately, one may administer the compound in a local rather than systemic manner, for example, via injection of the compound directly into a solid
5 tumor, often in a depot or sustained release formulation.

Furthermore, one may administer the drug in a targeted drug delivery system, for example, in a liposome coated with tumor-specific antibody. The liposomes will be targeted to and taken up selectively by the tumor.

10 **B. Composition/Formulation.**

Pharmacological compositions of the compounds and the pharmaceutically acceptable salts thereof are preferred embodiments of this invention. Pharmacological compositions of the present invention may be manufactured by processes well known in the art; e.g., by means of conventional
15 mixing, dissolving, granulating, dragee-making, levigating, emulsifying, encapsulating, entrapping or lyophilizing processes.

Pharmacological compositions for use in accordance with the present invention thus may be formulated in a conventional manner using one or more pharmaceutically acceptable carriers comprising excipients and auxiliaries which
20 facilitate processing of the active compounds into preparations which can be used pharmaceutically. Proper formulation is dependent upon the route of administration chosen.

For injection, the compounds of the invention may be formulated as sterile aqueous solutions, preferably in physiologically compatible buffers such as
25 Hanks' solution, Ringer's solution, or physiological saline buffer. For transmucosal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art.

For oral administration, the compounds can be formulated readily by combining the active compounds with pharmaceutically acceptable carriers well
30 known in the art. Such carriers enable the compounds of the invention to be formulated as tablets, pills, dragees, capsules, liquids, gels, syrups, slurries, suspensions and the like, for oral ingestion by a patient to be treated.

Pharmacological preparations for oral use can be made with the use of a solid excipient, optionally grinding the resulting mixture, and processing the mixture of granules, after adding suitable auxiliaries, if desired, to obtain tablets or dragee cores. Suitable excipients are, in particular, fillers such as sugars, including
5 lactose, sucrose, mannitol, or sorbitol; cellulose preparations such as, for example, maize starch, wheat starch, rice starch, potato starch, gelatin, gum tragacanth, methyl cellulose, hydroxypropylmethyl cellulose, sodium carboxymethylcellulose, and/or polyvinylpyrrolidone (PVP). If desired, disintegrating agents may be added, such as the cross-linked polyvinyl
10 pyrrolidone, agar, or alginic acid or a salt thereof such as sodium alginate.

Dragee cores are provided with suitable coatings. For this purpose, concentrated sugar solutions may be used, which may optionally contain gum arabic, talc, polyvinyl pyrrolidone, carbopol gel, polyethylene glycol, and/or titanium dioxide, lacquer solutions, and suitable organic solvents or solvent
15 mixtures. Dyestuffs or pigments may be added to the tablets or dragee coatings for identification or to characterize different combinations of active compound doses.

Pharmacological compositions which can be used orally include push-fit capsules made of gelatin, as well as soft, sealed capsules made of gelatin and a
20 plasticizer, such as glycerol or sorbitol. The push-fit capsules can contain the active ingredients in admixture with a filler such as lactose, binders such as starches, and/or lubricants such as talc or magnesium stearate and, optionally, stabilizers. In soft capsules, the active compounds may be dissolved or suspended in suitable liquids, such as fatty oils, liquid paraffin, or liquid
25 polyethylene glycols. In addition, stabilizers may be added. All formulations for oral administration should be in dosages suitable for such administration.

For buccal administration, the compositions may take the form of tablets or lozenges formulated in conventional manner.

For administration by inhalation, the compounds for use according to the
30 present invention are conveniently delivered in the form of an aerosol spray presentation from pressurized packs or a nebulizer, with the use of a suitable propellant, e.g., dichlorodifluoromethane, trichlorofluoromethane,

dichlorotetrafluoroethane, carbon dioxide or other suitable gas. In the case of a pressurized aerosol the dosage unit may be determined by providing a valve to deliver a metered amount. Capsules and cartridges of, e.g., gelatin for use in an inhaler or insufflator may be formulated containing a powder mix of the compound and a suitable powder base such as lactose or starch.

The compounds may be formulated for parenteral administration, e.g., by bolus injection or continuous infusion. Formulations for injection may be presented in unit dosage form, e.g., in ampules or in multi-dose containers, with an added preservative. The compositions may take such forms as suspensions, solutions or emulsions in oily or aqueous vehicles, and may contain formulatory agents such as suspending, stabilizing and/or dispersing agents.

Pharmacological compositions for parenteral administration include sterile aqueous solutions of the active compounds in water soluble form. Additionally, suspensions of the active compounds may be prepared as appropriate oily injection suspensions. Suitable lipophilic solvents or vehicles include fatty oils such as sesame oil, or synthetic fatty acid esters, such as ethyl oleate or triglycerides, or liposomes. Aqueous injection suspensions may contain substances which increase the viscosity of the suspension, such as sodium carboxymethyl cellulose, sorbitol, or dextran. Optionally, the suspension may also contain suitable stabilizers or agents which increase the solubility of the compounds to allow for the preparation of highly concentrated solutions.

Alternatively, the active ingredient may be in powder form for constitution with a suitable vehicle, e.g., sterile pyrogen-free water, before use.

The compounds may also be formulated in rectal compositions such as suppositories or retention enemas, e.g., containing conventional suppository bases such as cocoa butter or other glycerides.

In addition to the formulations described previously, the compounds may also be formulated as a depot preparation (*see*, for example, U.S. Patent No. 5,702,717 for a biodegradable depot for the delivery of a drug). Such long acting formulations may be administered by implantation (for example subcutaneously or intramuscularly) or by intramuscular injection. Thus, for example, the compounds may be formulated with suitable polymeric or hydrophobic materials

(for example as an emulsion in an acceptable oil) or ion exchange resins, or as sparingly soluble derivatives, for example, as a sparingly soluble salt. The pharmacological compositions herein also may comprise suitable solid or gel phase carriers or excipients. Examples of such carriers or excipients include but
5 are not limited to calcium carbonate, calcium phosphate, various sugars, starches, cellulose derivatives, gelatin, and polymers such as polyethylene glycols.

The compounds of the invention that inhibit LPAAT- β may be provided as physiologically acceptable salts wherein the claimed compound may form the negatively or the positively charged species. Examples of salts in which the
10 compound forms the positively charged moiety include, without limitation, quaternary ammonium (defined elsewhere herein), salts such as the hydrochloride, sulfate, carbonate, lactate, tartrate, maleate, succinate, etc. formed by the reaction of an amino group with the appropriate acid.

15 **C. Dosage.**

Pharmacological compositions suitable for use in the present invention include compositions wherein the active ingredients are contained in an amount effective to achieve its intended purpose.

More specifically, a therapeutically effective amount means an amount of
20 compound effective to prevent, alleviate or ameliorate symptoms of disease or prolong the survival of the subject being treated.

Determination of a therapeutically effective amount is well within the capability of those skilled in the art, especially in light of the detailed disclosure provided herein.

For any compound used in the methods of the invention, the
25 therapeutically effective amount or dose can be estimated initially from cell culture assays. For example, a dose can be formulated in animal models to achieve a circulating concentration range that includes the IC_{50} as determined in cell culture (i.e., the concentration of the test compound which achieves a half-
30 maximal inhibition of LPAAT- β activity). Such information can be used to more accurately determine useful doses in humans.

Toxicity and therapeutic efficacy of the compounds described herein can be determined by standard pharmaceutical procedures in cell cultures or experimental animals for determining the LD₅₀ (the dose lethal to 50% of the population) and the ED₅₀ (the dose therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index and it can be expressed as the ratio between LD₅₀ and ED₅₀. Compounds which exhibit high therapeutic indices are preferred. The data obtained from these cell culture assays and animal studies can be used in formulating a range of dosage for use in human. The dosage of such compounds lies preferably within a range of circulating concentrations that include the ED₅₀ with little or no toxicity. The dosage may vary within this range depending upon the dosage form employed and the route of administration utilized. The exact formulation, route of administration and dosage can be chosen by the individual physician in view of the patient's condition. (see e.g., Fingl, *et al.*, in "The Pharmacological Basis of Therapeutics," (1975), Chapter 1, pp. 1).

Dosage amount and interval may be adjusted individually to provide plasma levels of the active moiety which are sufficient to maintain LPAAT- β inhibitory effects, or minimal effective concentration (MEC). The MEC will vary for each compound but can be estimated from in vitro data; e.g., the concentration necessary to achieve 50-90% inhibition of LPAAT- β using the assays described herein. Dosages necessary to achieve the MEC will depend on individual characteristics and route of administration. However, HPLC assays or bioassays can be used to determine plasma concentrations.

Dosage intervals can also be determined using MEC value. Compounds should be administered using a regimen which maintains plasma levels above the MEC for 10-90% of the time, preferably between 30-90% and most preferably between 50-90%.

In cases of local administration or selective uptake, the effective local concentration of the drug may not be related to plasma concentration.

The amount of composition administered will, of course, be dependent on the subject being treated, on the subject's weight, the severity of the affliction, the manner of administration and the judgment of the prescribing physician. An

exemplary systemic daily dosage is about 5 to about 200 mg/kg of body weight. Normally, from about 10 to about 100 mg/kg of body weight of the compounds of the preferred embodiments of the present invention, in one or more dosages per day, is effective to obtain the desired results. One of ordinary skill in the art can
5 determine the optimal dosages and concentrations of the compounds of the preferred embodiments of the present invention with only routine experimentation.

The compounds of the preferred embodiments of the present invention are substantially pure and preferably sterile. The phrase "substantially pure"
10 encompasses compounds created by chemical synthesis and/or compounds substantially free of chemicals which may accompany the compounds in the natural state, as evidenced by thin layer chromatography (TLC) or high performance liquid chromatography (HPLC).

15 **D. Other Applications**

The compounds of the preferred embodiments of the present invention may be employed not only for therapeutic purposes, but also as aids in performing research *in vitro*. For example, the compounds of the preferred
20 embodiments of the present invention may be used to study biochemical pathways that would require the inhibition of LPAAT- β to elevated levels of LPA. Inhibition of LPAAT- β may result in the prolonged or limited activity of biochemical pathways that depend on, or respond to, elevated levels of LPA.

Additionally, a cell culture medium comprising the compounds of the preferred embodiments of the present invention is within the scope of the
25 invention.

V. Assays for LPAAT- β DNA, RNA and Protein.

DNA molecules encoding the human LPAAT- β gene, or fragments thereof, can be used to detect the level of LPAAT- β gene expression in tissue samples.

30 Such a detection method can be used, for example, to compare the amount of LPAAT- β RNA in a sample obtained from normal tissue and in a sample isolated from methotrexate-resistant tumor tissue. The presence of relatively low levels of

LPAAT- β RNA in the tumor sample would indicate that methotrexate resistance is due, at least in part, to underexpression of the LPAAT- β gene.

In testing a tissue sample for LPAAT- β RNA using a nucleic acid hybridization assay, RNA can be isolated from tissue by sectioning on a cryostat and lysing the sections with a detergent such as SDS and a chelating agent such as EDTA, optionally with overnight digestion with proteinase K. Such tissue may be obtained by biopsy. A preferred quantity of tissue is in the range of 10-100 milligrams. Protein may be removed by phenol and chloroform extractions, and nucleic acids are precipitated with ethanol. RNA may be isolated by chromatography on an oligo dT column and then eluted from the column. Further fractionation can also be carried out according to methods well known to those of ordinary skill in the art.

A number of techniques for molecular hybridization are used for the detection of DNA or RNA sequences in tissues. When large amounts of tissue are available, analysis of hybridization kinetics provides the opportunity to accurately quantitate the amount of DNA or RNA present, as well as to distinguish sequences that are closely related but not identical to the probe. Reactions are run under conditions of hybridization (T_m -25 °C) in which the rate of re-association of the probe is optimal. Wetmur *et al.*, *J. Mol. Biol.* 31:349 (1968). The kinetics of the reaction are second order when the sequences in the tissue are identical to those of the probe; however, the reaction exhibits complex kinetics when probe sequences have partial homology to those in the tissue. Sharp *et al.*, *J. Mol. Biol.* 86:709 (1974).

The concentration of probe to cellular RNA is determined by the sensitivity desired. To detect one transcript per cell would require about 100 pg of probe per mg of total cellular DNA or RNA. The nucleic acids are mixed, denatured, brought to the appropriate salt concentration and temperature, and allowed to hybridize for various periods of time. The rate of reassociation can be determined by quantitating the amount of probe hybridized either by hydroxyapatite chromatography (Britten *et al.*, *Science* 161:529 (1968)) or by S1 nuclease digestion (Sutton, *Biochim. Biophys. Acta* 240:522 (1971)).

Another method of hybridization is the Northern Blot technique. The

particular hybridization technique is not essential to the invention, and any technique commonly used in the art is within the scope of the present invention. Typical probe technology is described in U. S. Patent No. 4,358,535, incorporated by reference herein. For example, hybridization can be carried out in a solution
5 containing 6 x SSC (10 x SSC: 1.5 M sodium chloride, 0.15 M sodium citrate, pH 7.0), 5 x Denhardt's (1 x Denhardt's: 0.2% bovine serum albumin, 0.2% polyvinylpyrrolidone, 0.02% Ficoll 400), 10 mM EDTA, 0.5% SDS and about 10^7 cpm of nick-translated DNA for 16 hours at 65 °C.

The aforementioned hybridization assays are particularly well suited for
10 preparation and commercialization in kit form, the kit comprising a carrier means compartmentalized to receive one or more container means (vial, test tube, etc.) in close confinement, with each container means comprising one of the separate elements to be used in hybridization assay. For example, there may be a container means containing LPAAT- β DNA molecules suitable for labeling by "nick translation," or containing labeled LPAAT- β DNA or labeled LPAAT- β RNA
15 molecules. Further container means may contain standard solutions for nick translation of DNA comprising DNA polymerase I/DNase I and unlabeled deoxyribonucleotides.

Antibodies to human LPAAT- β protein can be obtained using the product
20 of an LPAAT- β expression vector as an antigen. The preparation of polyclonal antibodies is well-known to those of skill in the art. See, for example, Green *et al.*, "Production of Polyclonal Antisera," in *Immunochemical Protocols* (Manson, ed.), pp. 1-5 (Humana Press 1992). Alternatively, an LPAAT- β antibody of the present invention may be derived from a rodent monoclonal antibody (MAb). Rodent
25 monoclonal antibodies to specific antigens may be obtained by methods known to those skilled in the art. See, for example, Kohler and Milstein, *Nature* 256:495, 1975, and Coligan *et al.* (eds.), *Current Protocols in Immunology*, 1:2.5.1-2.6.7 (John Wiley & Sons 1991) [hereinafter "Coligan"]. Briefly, monoclonal antibodies can be obtained by injecting mice with a composition comprising an antigen,
30 verifying the presence of antibody production by removing a serum sample, removing the spleen to obtain B-lymphocytes, fusing the B-lymphocytes with myeloma cells to produce hybridomas, cloning the hybridomas, selecting positive

clones which produce antibodies to the antigen, culturing the clones that produce antibodies to the antigen, and isolating the antibodies from the hybridoma cultures.

MAbs can be isolated and purified from hybridoma cultures by a variety of techniques that are well known in the art. Such isolation techniques include affinity
5 chromatography with Protein-A Sepharose, size-exclusion chromatography, and ion-exchange chromatography. See, for example, Coligan at pages 2.7.1-2.7.12 and pages 2.9.1-2.9.3. Also, see Baines *et al.*, "Purification of Immunoglobulin G (IgG)," in *Methods in Molecular Biology*, 10:79 (Humana Press, Inc. 1992). A LPAAT- β antibody may also be derived from a subhuman primate antibody.
10 General techniques for raising therapeutically useful antibodies in baboons may be found, for example, in Goldenberg *et al.*, International Patent Publication No. WO 91/11465 (1991), and in Losman *et al.*, *Int. J. Cancer* 46:310 (1990).

Alternatively, a therapeutically useful LPAAT- β antibody may be derived from a "humanized" monoclonal antibody. Humanized monoclonal antibodies are
15 produced by transferring mouse complementary determining regions from heavy and light variable chains of the mouse immunoglobulin into a human variable domain, and then, substituting human residues in the framework regions of the murine counterparts. The use of antibody components derived from humanized monoclonal antibodies obviates potential problems associated with the
20 immunogenicity of murine constant regions. General techniques for cloning murine immunoglobulin variable domains are described, for example, by the publication of Orlandi *et al.*, *Proc. Nat'l. Acad. Sci. USA* 86:3833 (1989). Techniques for producing humanized MAbs are described, for example, by Jones *et al.*, *Nature* 321:522 (1986); Riechmann *et al.*, *Nature* 332:323 (1988); Verhoeyen *et al.*,
25 *Science* 239:1534 (1988); Carter *et al.*, *Proc. Nat'l Acad. Sci. USA* 89:4285 (1992); Sandhu, *Crit. Rev. Biotech.* 12: 437 (1992); and Singer *et al.*, *J. Immun.* 150:2844 (1993), each of which is hereby incorporated by reference.

As an alternative, a LPAAT- β antibody of the present invention may be derived from human antibody fragments isolated from a combinatorial
30 immunoglobulin library. See, for example, Barbas *et al.*, *METHODS: A Companion to Methods in Enzymology* 2:119 (1991); and Winter *et al.*, *Ann. Rev. Immunol.* 12:433 (1994) which are incorporated herein by reference. Cloning and

expression vectors that are useful for producing a human immunoglobulin phage library can be obtained, for example, from STRATAGENE Cloning Systems (La Jolla, CA). In addition, a LPAAT- β antibody of the present invention may be derived from a human monoclonal antibody. Such antibodies are obtained from
5 transgenic mice that have been "engineered" to produce specific human antibodies in response to antigenic challenge. In this technique, elements of the human heavy and light chain locus are introduced into strains of mice derived from embryonic stem cell lines that contain targeted disruptions of the endogenous heavy chain and light chain loci. The transgenic mice can synthesize human antibodies specific for
10 human antigens, and the mice can be used to produce human antibody-secreting hybridomas. Methods for obtaining human antibodies from transgenic mice are described by Green *et al.*, *Nature Genet.* 7:13 (1994); Lonberg *et al.*, *Nature* 368:856 (1994); and Taylor *et al.*, *Int. Immun.* 6:579 (1994).

Having now generally described this invention, the same will be
15 understood by reference to the following examples which are provided herein for purposes of illustration only and are not intended to be limiting unless otherwise specified.

Example 1

Production of recombinant LPAAT- β for Various Assays

For the construction of Baculovirus expression vectors, the full-length human LPAAT- β cDNA was amplified by PCR from the DNA template pCE9.LPAAT- β (West *et al.*, *DNA Cell Biol.* 16:691-701 (1997)) using the
25 primers 5'-TGATATCCGA AGAAGATCTT ATGGAGCTGT GGCCGTGTC-3' (olpb1F) and 5'-CAGGCTCTAG ACTACTGGGC CGGCTGCAC-3' (olpb1R). The ~870 bp fragment generated was reamplified by PCR using the primers 5' CCTACGTCG ACATGGAACA AAAATTGATA TCCGAAGAAG ATC-3' (olpb2F) and 5'-CAGGCTCTAG ACTACTGGGC CGGCTGCAC-3'
30 (olpb1R). The ~890 bp fragment generated was then cleaved with Sal I and Xba I for insertion into pFastBac™ HTc vector (Life Technologies, Gaithersburg, MD) between the Sal I and Xba I sites for the generation of the plasmid pFB.LPAAT- β . This plasmid was then transformed into *E. coli* DH10Bac™ (Life

Technologies, Gaithersberg, MD) for the generation of recombinant Bacmid DNA for transfection into HighFive (Invitrogen, San Diego, CA) or Sf9 insect cells for the production of recombinant Baculovirus stocks using the protocol described in the Bac-to-Bac® Baculovirus Expression System (Life Technologies, Gaithersberg, MD), a eukaryotic expression system for generating recombinant baculovirus through site-specific transposition in *E. coli*. Viral stocks harvested from the transfected cells can then be used to infect fresh insect cells for the subsequent expression of LPAAT- β fusion protein with a poly-histidine tag and a myc-epitope near its N-terminus. The membrane fraction from these Sf9 cells would be the source of LPAAT enzyme.

Example 2

Preparation of cell membranes from Sf9 cells

For the preparation of membranes From Sf9 Cells, all steps are performed on ice or at 4 °C. Sf9 cell pellets ($\sim 10^8$ cells) were thawed and resuspended in 1-2 ml of buffer A (20 mM Hepes, pH 7.5, 1 mM DTT, 1 mM EDTA, 20% w/v glycerol, 1 mM Benzamidine, 1 μ g/ml soybean trypsin inhibitor (SBTI), 1 μ g/ml pepstatin A) w/o DTT but with 1 mM Pefabloc. The cells were lysed by sonication using a Branson Sonifier at output = 2, duty cycle = 2, 10 pulses each at 10s. with the tip of small sonicator probe submerged but not touching the walls. DTT was then added to 1 mM from a 1 M stock. The samples were centrifuged at 1500 rpm for 5 min. The low speed supernatant was saved and centrifuged (TLA 100.3 rotor, polycarbonate tubes, 2 ml/tube or 1.5 ml/tube minimum) at 100000 x g for 1 hr. The high speed pellet was resuspend in Buffer A with a probe sonicator (10 pulses @ output #2 and duty cycle 20%) as a source of LPAAT enzyme.

Example 3

Assay of LPAAT- β Activity

LPAAT- β catalyzes the transfer of an acyl group from a donor such as acyl-CoA to LPA. The transfer of the acyl group from acyl-CoA to LPA leads to

the release of free CoA, which can be reacted with the thiol reagent, 5,5'-dithiobis(2-nitrobenzoic acid) (DTNB). The reaction between DTNB and the free
sulfhydryl group from CoA generates a yellow-colored product, 3-carboxylato-4-nitrothiophenolate (CNP), that absorbs at 413 nm. LPAAT- β derived from Sf9
5 cell membrane overexpressing LPAAT- β were resuspended in HEPES saline
buffer (20 mM HEPES pH 7.5, 150 mM NaCl), 1 mg/ml BSA and 72 μ l aliquots
were distributed into 96-well microtiter plates. 8 μ l of compound of interest at
200 μ M dissolved in 100% DMSO was added into each well. 20 μ l of 1 mM
18:1-CoA and 1 mM sn-1-18:1 lysoPA was then added to each well to initiate the
10 reaction and allowed to run at room temperature for 25 min. 100 μ l of 1 mM
DTNB in 100% ethanol was then added to each well to quench the reaction and
for color development. The absorbance at 405 nm, measured using a
spectrophotometer plate reader, is proportional to the activity of LPAAT- β in the
sample. This colorimetric assay was used for the high throughput screening of
15 LPAAT inhibitors. Compounds that showed >50% inhibition of the change in
absorbance at 405 nm compared to control were selected for a secondary assay.
Figure 7 shows an example of the colorimetric assay of which the time course of
color development is dependent on the amount of LPAAT enzyme added. Figure
8 shows an example of the results obtained from assaying a plate of various
20 compounds at 16 μ M. Compounds that gave a reading of less than 0.06 arbitrary
units (indicated by arrow on right margin) were selected for further study.

A secondary assay for LPAAT activity in cell extracts based on either the
conversion of fluorescent NBD-LPA to NBD-PA (West, et al., *DNA Cell Biol.* 6:
691-701, 1997) or [14 C]LPA to [14 C]PA using TLC analysis was used to screen
25 compounds that showed >50% inhibition of LPAAT activity in the primary
colorimetric assay. The radiometric assay was carried out in Sf9 cell membrane
overexpressing LPAAT- β resuspended in HEPES-saline buffer, pH 7.5, 1 mg/ml
BSA, 1mM EDTA and 200 μ M [14 C]18:1-CoA and 200 μ M sn-1-18:1 lysoPA.
The samples were incubated 7 min at 37 $^{\circ}$ C, extracted into organic solvent
30 (CHCl_3 / CH_3OH / HCl at 33/66/1), before loading onto TLC plates. A more
detailed protocol for the radiometric assay is described below:

Specifically, this LPAAT assay is a modification of the acyltransferase

assay published previously (Hollenback and Glomset, *Biochemistry* 37: 363-376 (1999)).

1. The basic assay, in a total vol of 50 μ l, employs a solution of substrates and the protein sample. Total assay volume, as well as the volume of
5 each solution, can be changed to fit an experiment. In addition, other compounds, ex inhibitors and activators, can be included in the assay as well.

2. To prepare the solution of substrates:

a. Stocks of Hepes (pH 7.5), NaCl, EDTA, BSA and acyl-CoA (from Serdery or Sigma) are mixed with water to make the appropriate concentration of
10 each compound. This can be varied from assay-to-assay, but the final reaction mix is about 50 mM Hepes, 100 mM NaCl, 1 mM EDTA, 1 mg/ml BSA and 0-400 μ M acyl-CoA.

b. The lysoPA (from Avanti) is typically stored in chloroform and the 14 C-labeled acyl-CoA (from Amersham) is typically stored in
15 water/ethanol=1:1. Appropriate amounts of each solution are added the to a 12 x 75 mm borosilicate glass test tube and dry the solvent under N_2 or Ar. An appropriate volume of the solution prepared in 2a is added to the lysoPA and 14 C-labeled acyl-CoA. The lipids are resuspend by sonication for 15 sec in a bath sonicator. The resulting suspension is then incubated (with occasional gentle
20 vortexing) for about 10 minutes at room temp. The *sn*-1-16:0 lysoPA may require brief warming of the solvent to solubilize it. The concentration of lysoPA and 14 C-labeled acyl-CoA can vary, but typically the final lysoPA concentration ranges between 0 and 400 μ M and the 14 C-labeled acyl-CoA specific activity ranges between 0.5 and 2 Ci/mol.

25 3. Protein sample: varies from experiment-to-experiment.

4. The assay is performed by mixing the components in 12 x 75 mm borosilicate glass test tubes (the order of addition does not matter unless indicated) and incubating at 37 $^{\circ}$ C for 5 to 10 minutes such that the assay within
the linear range for time and protein.

30 5. The reaction is quenched by adding 1.3 ml of chloroform/methanol/HCl = 48/51/0.7 and vortexing. 10 μ l of carrier solution is then added (3 mg/ml each PA, ex. 16:0-18:1, and lysoPA, ex *sn*-1-18:1, in

chloroform). Two phases are formed by adding 0.3 ml of water to each tube and vortexing.

6. The sample is centrifuged for 3 minutes at 1000 x g, the upper (aqueous/methanol) phase is aspirated and the lower phase is dried under
5 nitrogen.

7. Thin layer chromatography:

a. The dried samples are resuspended in 50 μ l of chloroform and a 15 μ l aliquot is immediately spotted on an Analtech silica gel 60 HP-TLC plate (10 x 20 cm).

10 b. Plates are developed in chloroform/methanol/acetic acid/water = 85/12.5/12.5/3 (takes about 15 min) and dried.

c. To be able to convert pixel volume (determined by the Storm phosphor imager, see step 8b) into cpm, cpm standard curve must be generated on the plate. 14 C-labeled oleate dilutions in chloroform are made for this purpose.
15 Four stocks (50 cpm/ μ l to 800 cpm/ μ l) are made and 2 μ l of a different concentration are spotted in each corner of the plate (where previously there was no radioactivity).

d. For quality control purposes, the plates are stained with primuline and scanned with the Storm (blue chemilluminescence mode).

20 The PA and lysoPA bands are easily detected in this system because of the carrier added in step 5. PA and lysoPA have respective R_f's of about 0.63 and 0.21.

8. Quantitating activity:

a. The plates are then wrapped in saran wrap and exposed to a freshly
25 blanked phosphor screen overnight (longer exposures can also be done to increase the signal).

b. The screens are scanned (Phosphorimager mode), and LPAAT activity is determined by quantifying the pixels in the band comigrating with PA standard versus the standard curve generated from the cpm standards that were
30 spotted in step 7c.

Figure 9 shows examples of a compound, namely, 6-chloro-N,N'-diphenyl-[1,3,5]triazine-2,4-diamine, selected from the secondary screening that

exhibit concentration-dependent inhibition of LPAAT- β activity (O). Moreover, these compounds have minimal effect on LPAAT- α activity (), suggesting they are isoform-specific inhibitors.

5

Example 4

Triazines of the preferred embodiments of the present invention were synthesized by one of two methods. Symmetrically substituted triazines were synthesized by the addition of two equivalents of the appropriate amino
10 compound, in the presence of diisopropylethylamine, to cyanuric chloride. Non-symmetrical triazines were synthesized in a stepwise fashion by the sequential addition of the amino compound in the presence of potassium carbonate with isolation of the intermediate mono-amino-dichlorotriazine.

15 A. Method 1: Synthesis of Symmetrical Triazines.

Synthesis of 6-Chloro-N,N'-diphenyl-[1,3,5]triazine-2,4-diamine (CT-113020)

To a mixture of cyanuric chloride (5.07 g, 27.5 mmol) and acetonitrile
20 (60 ml), cooled in an ice bath, was added a solution of aniline (5.3 ml, 58.2 mmol) and diisopropylethylamine (10.5 ml, 60.3 mmol) in acetonitrile (15 ml) over 30 minutes. After stirring at room temperature for 20 hours, the mixture was filtered. The solid was washed with ethyl acetate (4x 25 ml), suspended in water (50 ml), stirred for 1 hour, filtered, washed with water, and dried
25 under vacuum to give 6-chloro-N,N'-diphenyl-[1,3,5]triazine-2,4-diamine (3.97 g, 48% yield) as a white solid. ¹H NMR (*d*₆-DMSO) δ 7.61 (2 H, br s), 7.28-7.37 (4 H, m), 7.05-7.11 (4 H, m). ¹³C NMR (*d*₆-DMSO) δ 167.7, 164.0, 138.5, 128.4, 124.5, 121.3.

30 B. Method 2: Synthesis of Unsymmetrical Triazines.

Synthesis of 6-Chloro-N-(4-chlorophenyl)-N'-phenyl-[1,3,5]triazine-2,4-diamine (CT-116433)

To a mixture of cyanuric chloride (5.15 g, 27.9 mmol), potassium

carbonate (3.98 g, 28.8 mmol) and 18-crown-6 (158 mg, 0.60 mmol) in toluene (40 ml), cooled in an ice bath, was added a solution of 4-chloroaniline (3.61 g, 28.3 mmol) in toluene (20 ml) over 15 minutes. After stirring at room temperature for 20 hours, the mixture was treated with ethyl acetate (60 ml) and filtered through a pad of celite under suction. The filtrate was concentrated under vacuum and recrystallized (chloroform) to give (4-chloro-phenyl)-(4,6-dichloro-[1,3,5]triazin-2-yl)-amine (4.06 g, 53% yield) as a white solid. ¹H NMR (*d*₆-DMSO) δ 11.23 (1 H, s), 7.62 (2 H, d, *J* = 9 Hz), 7.47 (2 H, d, *J* = 9 Hz). ¹³C NMR (*d*₆-DMSO) δ 169.5, 168.5, 163.6, 135.7, 128.6, 122.5, ESMS *m/z* 273 (*M* - H)⁺.

To a mixture of (4-chloro-phenyl)-(4,6-dichloro-[1,3,5]triazin-2-yl)-amine (3.97 g, 14.4 mmol), potassium carbonate (2.20 g, 15.9 mmol) and 18-crown-6 (46 mg, 0.17 mmol) in toluene (25 ml), cooled in an ice bath, was added a solution of aniline (1.4 ml, 15.4 mmol) in toluene (10 ml) over 15 minutes. After stirring at room temperature for 24 hours, the mixture was treated with ethyl acetate (35 ml) and filtered through a pad of celite under suction. The filtrate was concentrated under vacuum and the residue was recrystallized (chloroform) to give 6-Chloro-N-(4-chlorophenyl)-N'-phenyl-[1,3,5]triazine-2,4-diamine (2.20 g, 46% yield) as a white solid. ¹H NMR (*d*₆-DMSO) δ 10.1-10.4 (2 H, br s), 7.5-7.8 (4 H, br s), 7.3-7.5 (4 H, m), 7.15-7.05 (1 H, m), ESMS *m/z* 330 (*M* - H)⁺.

Example 5

Synthesis of 6-Chloro-N-(4-chlorophenyl)-N'-(4-methoxyphenyl)-[1,3,5]triazine-2,4-diamine (CT-31867)

The reaction of (4-chloro-phenyl)-(4,6-dichloro-[1,3,5]triazin-2-yl)-amine with *p*-anisidine, according to method 2, gave 6-chloro-N-(4-chlorophenyl)-N'-(4-methoxyphenyl)-[1,3,5]triazine-2,4-diamine (51 mg, 62%): ¹H NMR (CDCl₃) 7.21-7.50 (6 H, m), 6.91 (2 H, d, *J* = 11 Hz), 3.85 (3 H, s).

Example 6

Synthesis of 6-Chloro-N-(4-methoxyphenyl)-N'-phenyl-[1,3,5]triazine-2,4-diamine (CT-31942)

5 To a solution of 2-(4-methoxyphenyl)amino-4,6-dichloro-1,3,5-triazine (57 mg, 0.21 mmoles) in acetonitrile (0.5 ml) was added a solution of aniline (0.021 ml, 0.23 mmoles) and diisopropylethylamine (0.05 ml, 0.29 mmoles) in acetonitrile (0.5 ml). After stirring for 24 hours, the mixture was concentrated under vacuum and the residue was dissolved in ethyl acetate (10 ml). The
10 solution was washed with a solution composed of saturated aqueous sodium chloride solution and 1 M hydrochloric acid (1:1, 2 x 10 ml), dried over sodium sulfate, filtered, and concentrated under vacuum. The residue was purified by flash chromatography on silica gel eluting with 10% ethyl acetate-hexane to provide CT-31942 (38 mg, 57% yield). ¹H NMR (*d*₆-DMSO) δ
15 10.00-10.22 (m, 2 H), 7.26-7.82 (m, 6 H), 7.01-7.10 (m, 1 H), 6.91 (d, *J* = 9 Hz, 2 H), 3.74 (s, 3 H).

Example 7

Synthesis of N-Benzo[1,3]dioxol-5-yl-6-chloro-N'-(4-chlorophenyl)-[1,3,5]triazine-2,4-diamine (CT-31978)

20 The reaction of 4,6-dichloro-N-(4-chlorophenyl)-[1,3,5]triazine-2-amine with 3,4-methylenedioxyaniline using the method described for the synthesis of CT-116433 provided CT-31978 (56 mg, 65% yield). ¹H NMR (*d*₆-DMSO) δ 10.08-10.37 (m, 2 H), 7.58-7.87 (m, 2 H), 7.21-7.40 (m, 3 H),
25 6.85-7.03 (m, 2 H), 6.00 (s, 2 H).

Example 8

Synthesis of 6-Chloro-N-(2,3-dihydrobenzo[1,4]dioxin-6-yl)-N'-phenyl-[1,3,5]triazine-2,4-diamine (CT-32028)

30 To a solution of 4,6-dichloro-N-phenyl-[1,3,5]triazine-2-amine (95 mg, 0.39 mmoles) in tetrahydrofuran (2 ml) was added a solution of 1,4-benzodioxan-6-amine (64 mg, 0.42 mmoles) and triethylamine (0.07 ml, 0.50 mmoles) in tetrahydrofuran (0.5 ml). After stirring for 24 hours, the mixture was concentrated under vacuum and the residue was dissolved in ethyl acetate

(10 ml). The solution was washed with a solution composed of saturated aqueous sodium chloride solution and 1 M hydrochloric acid (1:1, 2x 10 ml), dried over sodium sulfate, filtered, and concentrated under vacuum. The residue was purified by flash chromatography on silica gel eluting with 25% ethyl acetate-hexanes to provide CT-32028 (97 mg, 69% yield). ¹H NMR (*d*₆-DMSO) δ 9.88-10.28 (m, 2 H), 7.60-7.85 (m, 2 H), 7.20-7.40 (m, 3 H), 6.96-7.10 (m, 2 H), 6.77-6.81 (m, 1 H), 4.21 (s, 4 H).

Example 9

Synthesis of *N*-Benzo[1,3]dioxol-5-yl-6-chloro-*N'*-phenyl-[1,3,5]triazine-2,4-diamine (CT-32042)

The reaction of 4,6-dichloro-*N*-phenyl-[1,3,5]triazine-2-amine with 3,4-methylenedioxyaniline using the method described for the synthesis of CT-32028 provided CT-32042 (51 mg, 42% yield). ¹H NMR (*d*₆-DMSO) δ 10.05-10.29 (m, 2 H), 7.57-7.80 (m, 2 H), 7.21-7.42 (m, 3 H), 6.80-7.13 (m, 3 H), 6.03 (s, 2 H).

Example 10

Synthesis of 6-Chloro-*N*-indan-5-yl-*N'*-phenyl-[1,3,5]triazine-2,4-diamine (CT-32099)

The reaction of 4,6-dichloro-*N*-phenyl-[1,3,5]triazine-2-amine with 5-aminoindan using the method described for the synthesis of CT-32028 provided CT-32099 (36 mg, 37% yield). ¹H NMR (*d*₆-DMSO) δ 10.15-10.28 (m, 2 H), 7.56-7.72 (m, 3 H), 7.05-7.39 (m, 5 H), 2.75-3.00 (m, 4 H), 1.94-2.09 (m, 2 H).

Example 11

Synthesis of 6-chloro-*N*-2-(4-chlorophenyl)-*N*-4-propyl-[1,3,5]triazine-2,4-diamine (CT-116988)

The reaction of 4,6-dichloro-*N*-(4-chlorophenyl)-[1,3,5]triazine-2-amine with propylamine using the method described for the synthesis of CT-116433 provides CT-116988.

Example 12

Synthesis of N-(4-Chlorophenyl)-6-methoxy-N'-propyl-[1,3,5]triazine-2,4-diamine (CT-117147)

5 A mixture of CT-116988 and sodium methoxide (3.0 molar equivalents) in methanol is heated at reflux for 18 hours. After cooling to room temperature, the reaction mixture is concentrated under vacuum. The residue is suspended in ethyl acetate and washed with water. The ethyl acetate phase is dried over sodium sulfate, filtered and concentrated under vacuum.
10 The residue is purified by flash chromatography on silica gel to provide CT-117147.

Example 13

Synthesis of N-(4-chlorophenyl)-6-methylsulfanyl-N'-phenyl-[1,3,5]triazine-2,4-diamine (CT-31888)

15 A mixture of CT-116433 (108 mg, 0.32 mmol) and sodium methanethiolate (79 mg, 1.13 mmol) in dimethyl sulfoxide (3 ml) was heated at 70 °C for 18 hours. After cooling to room temperature the mixture was diluted with ethyl acetate (25 ml) and was washed with a 1:1 solution of water
20 and saturated aqueous sodium chloride solution (4x 25 ml). The organic phase was dried over sodium sulfate, filtered, and concentrated under vacuum. The residue was purified by flash chromatography on silica gel eluting with 5% ethyl acetate-hexane to provide CT-31888 (76 mg, 69% yield). ¹H NMR (*d*₆-DMSO) δ 7.51-7.60 (m, 4 H), 7.26-7.39 (m, 5 H), 7.00-7.17 (m, 2 H), 2.56 (s,
25 3 H).

From the foregoing description, one skilled in the art can easily ascertain the essential characteristics of this invention, and without departing from the spirit and scope thereof, can make various changes and modifications of the invention to adapt it to various usages and conditions without undue
30 experimentation. All patents, patent applications and publications cited herein are incorporated by reference in their entirety.